

K080767

510(k) Summary
CONQUEST® Spinal System

MAY 21 2008

Submitted By: Life Spine
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510(k) Contact: Rebecca Brooks
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Date Prepared: March 14, 2008

Trade Name: CONQUEST® Spinal System

Common Name: Appliance, Fixation, Spinal Interlaminar

Classification: 888.3070 Pedicle screw spinal system

Device Product Code: MNH, 21 CFR 888.3070, Class II
MNI, 21 CFR 888.3070, Class II

Predicate Devices: ARX Spinal System (K061600)
Life Spine Cross Connector (K073430)

Device Description:

The CONQUEST Spinal System is a titanium alloy multiple component system comprised of a variety of non-sterile, single use implantable components. When assembled, the components create a rigid structure providing stabilization and promote spinal fusion. The system consists of an assortment of rods, screws, and locking caps.

Intended Use of the Device:

Internal fixation implants are load-sharing devices intended to stabilize and maintain alignment until normal healing occurs. Implants are not intended to replace normal body structures or bear the weight of the body in the presence of incomplete bone healing.

The CONQUEST Spinal System, when properly used, is intended for posterior pedicle screw fixation of the non-cervical posterior spine in skeletally mature patients. It provides stabilization and immobilization of spinal segments as an adjunct to fusion.

When used as a posterior spine thoracic/lumbar system, the CONQUEST Spinal System is indicated for one or more of the following: (1) degenerative disc disease (is defined as back pain of discogenic origin with degeneration of the disc confirmed by history and radiographic studies), (2) trauma (i.e. fracture or dislocation), (3) curvatures (scoliosis, kyphosis, and/or lordosis), (4) spinal tumor, (5) failed previous fusion (6) pseudarthrosis, (7) spinal stenosis, (8) spondylolisthesis.

Material:

Manufactured from medical grade titanium alloy described by ASTM F136 (Ti 6AL-4V-ELI) implant grade titanium alloy.

Performance Data:

Biomechanical testing in accordance with ASTM F1717 was conducted to demonstrate substantial equivalence.

Substantial Equivalence:

The CONQUEST® Spinal System was shown to be substantially equivalent to previously cleared devices in indications for use, design, function, and materials used.



Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Life Spine
% Ms. Rebecca Brooks
Project Manager
2401 W. Hassell Road, Suite 1535
Hoffman Estates, IL 60169

MAY 21 2008

Re: K080767
Trade/Device Name: Conquest Spinal System
Regulation Number: 21 CFR 888.3070
Regulation Name: Pedicle Screw Spinal System
Regulatory Class: II
Product Code: MNI, MNH
Dated: April 30, 2008
Received: May 6, 2008

Dear Ms. Brooks:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Center for Devices and Radiological Health's (CDRH's) Office of Compliance at (240) 276-0120. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding postmarket surveillance, please contact CDRH's Office of Surveillance and Biometric's (OSB's) Division of Postmarket Surveillance at 240-276-3474. For questions regarding the reporting of device adverse events (Medical Device Reporting (MDR)), please contact the Division of Surveillance Systems at 240-276-3464. You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at toll-free number (800) 638-2041 or (240) 276-3150 or the Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,

A handwritten signature in black ink, appearing to read "Mark N. Melkerson", with a stylized flourish at the end.

Mark N. Melkerson
Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indications for Use

510(k) number (if known): K080767

Device Name: CONQUEST® Spinal System

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Prescription Use x
(Part 21 CFR 801 Subpart D)

And/Or

Over-the-Counter Use
(21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF
NEEDED)

Neil Reed
(Division Sign-Off)

Division of General, Restorative,
and Neurological Devices

510(k) Number K080767